

**Technical report:** Regulation of DNA metabolism by DNA-binding proteins probed by single molecule spectroscopy

## A. Physiological background of DNA denaturation

The Watson-Crick double-helix is the thermodynamically stable configuration of a DNA molecule under physiological conditions (normal salt and room/body temperature). This stability is effected (a) by Watson-Crick H-bonding, that is essential for the specificity of base pairing, i.e., for the key-lock principle according to which the nucleotide Adenine exclusively binds to Thymine, and Guanine only to Cytosine. Base-pairing therefore guarantees the high level of fidelity during replication and transcription. (b) The second contribution to DNA helix stability comes from base-stacking between neighboring base pairs: through hydrophobic interactions between the planar aromatic bases, that overlap geometrically and electronically, the base-pair stacking stabilizes the helical structure against the repulsive electrostatic force between the negatively charged phosphate groups located at the outside of the DNA double-strand. While hydrogen bonds contribute only little to the helix stability, the major support comes from base-stacking.

The capability of the two complementary strands of DNA to move apart and to rejoin, without damaging the chemical structure of the two single-strands, is crucial to many physiological processes such as replication via the proteins DNA helicase and polymerase, and transcription through RNA polymerase. During these processes, the proteins unzip a certain region of the double-strand, to obtain access to the genetic information stored in the bases in the core of the double-helix. This unzipping corresponds to breaking the hydrogen bonds between the base-pairs. Classically, the so-called melting and reannealing behavior of DNA has been studied in solution in vitro by increasing the temperature, or by titration with acid or alkali (**Figure 1**). During thermal melting, the stability of the DNA duplex is related to the content of triple hydrogen-bonded G-C base-pairs: the larger the fraction of G-C pairs, the higher the required melting temperature or pH value. Thus, under thermal melting, dsDNA starts to unwind in regions rich in A-T base-pairs, and then proceeds to regions of progressively higher GC content. Conversely, molten, complementary chains of single-stranded DNA (ssDNA) begin to reassociate and eventually reform the original double-helix under incubation at roughly  $25^\circ$  below the melting temperature  $T_m$ .

Complementary to thermal or pH induced denaturation, ssDNA can be driven toward denaturation mechanically, by applying a tensional stress along the DNA in an optical tweezers trap. The force per extension increases in worm-like chain fashion, until a plateau at approximately 65 pN is reached (**Figure 2**). This plateau is sometimes interpreted as new DNA configuration, the S form. By a series of experiments, it appears more likely that the plateau corresponds to the mechanical denaturation transition. To first order, the effect of the longitudinal pulling with a force  $F$  translates into an external torque  $\tau = \tau(F)$ , whose effect is to decrease the free energy of melting  $\Delta G$ :

$$\Delta G(F) = \Delta G(F=0) - \tau(F) \theta_0 \quad (1)$$

where  $\theta_0 = 2\pi/10.35$  is the twist angle per bp of the double helix without applied forces.

20061226021

# REPORT DOCUMENTATION PAGE

Form Approved  
OMB No. 074-0188

Public reporting burden for this collection of information is estimated to average 1 hour per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing this collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to Washington Headquarters Services, Directorate for Information Operations and Reports, 1215 Jefferson Davis Highway, Suite 1204, Arlington, VA 22202-4302, and to the Office of Management and Budget, Paperwork Reduction Project (0704-0188), Washington, DC 20503

|                                  |                                |   |              |
|----------------------------------|--------------------------------|---|--------------|
| 1. AGENCY USE ONLY (Leave blank) | 2. REPORT DATE<br>Dec. 5, 2006 | 3. REPORT TYPE AND DATES COVERED<br>June 1, 2004-May 31, 2006 | <i>Final</i> |
|----------------------------------|--------------------------------|---|--------------|

|  |  |
|--|--|
| 4. TITLE AND SUBTITLE<br>Regulation of DNA metabolism by DNA-binding proteins probed by single molecule spectroscopy | 5. FUNDING NUMBERS<br>FA9550-04-1-0394 |
|--|--|

|                               |  |
|-------------------------------|--|
| 6. AUTHOR(S)<br>Andreas Hanke |  |
|-------------------------------|--|

|   |   |
|---|---|
| 7. PERFORMING ORGANIZATION NAME(S) AND ADDRESS(ES)<br>University of Texas at Brownsville /<br>Texas Southmost College<br>80 Fort Brown<br>Brownsville, TX 78520 | 8. PERFORMING ORGANIZATION<br>REPORT NUMBER |
|---|---|

|  |  |
|--|--|
| 9. SPONSORING / MONITORING AGENCY NAME(S) AND ADDRESS(ES)<br>AFOSR 875 N. Randolph St., Ste. 325,<br>RM 3112<br>Arlington, VA 22203-1768<br><i>Dr. Harold Weinstock/NE</i> | 10. SPONSORING / MONITORING<br>AGENCY REPORT NUMBER<br><br>AFRL-SR-AR-TR-06-0488 |
|--|--|

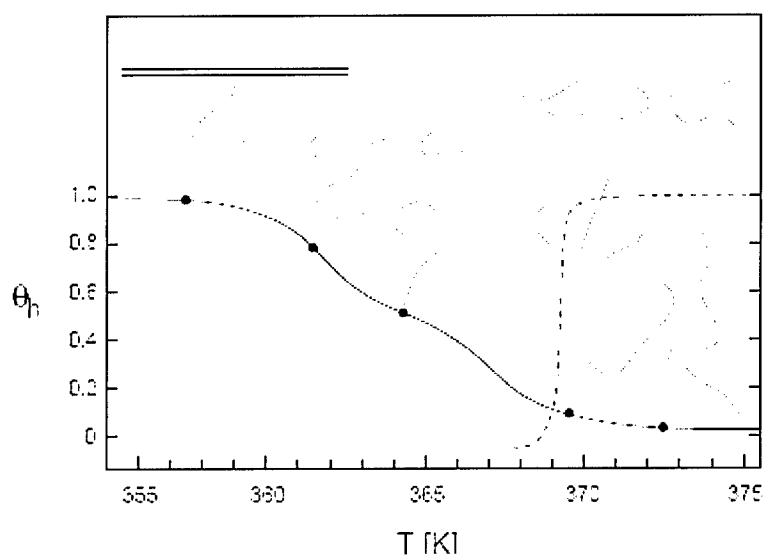
|                         |
|-------------------------|
| 11. SUPPLEMENTARY NOTES |
|-------------------------|

|   |                        |
|---|------------------------|
| 12a. DISTRIBUTION / AVAILABILITY STATEMENT<br>Approved for Public Release; distribution is unlimited. <i>A:</i> | 12b. DISTRIBUTION CODE |
|---|------------------------|

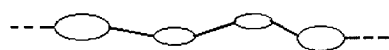
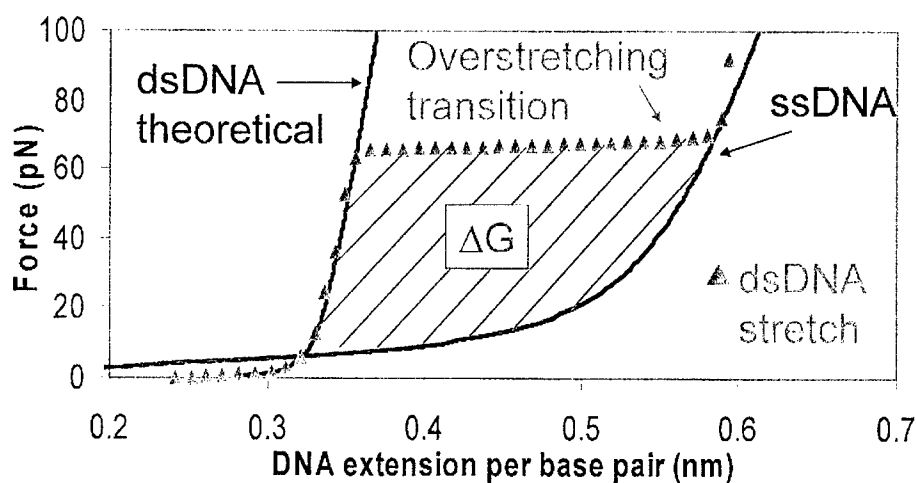
|   |
|---|
| 13. ABSTRACT (Maximum 200 Words)<br>Recent advances in single-molecule force spectroscopy of DNA make it possible to study the thermodynamics and kinetics of DNA binding proteins under a wide range of conditions. A biophysical model for the DNA binding T4 gene 32 protein has been developed to study the kinetics of DNA protein binding to transient single-stranded DNA regions due to thermal fluctuations. The model is used to analyze recent single-molecule spectroscopy data of this system. |
|---|

|                   |                          |
|-------------------|--------------------------|
| 14. SUBJECT TERMS | 15. NUMBER OF PAGES<br>8 |
|                   | 16. PRICE CODE           |

|  |  |  |                            |
|--|--|--|----------------------------|
| 17. SECURITY CLASSIFICATION<br>OF REPORT | 18. SECURITY<br>CLASSIFICATION<br>OF THIS PAGE | 19. SECURITY CLASSIFICATION<br>OF ABSTRACT<br>unclassified | 20. LIMITATION OF ABSTRACT |
|--|--|--|----------------------------|



**Figure 1.** Fraction  $\theta_h$  of double-helical domains within the DNA as a function of temperature  $T$ . Schematic representation of  $\theta_h(T)$ , showing the increased formation of bubbles and unzipping from the ends until full denaturation has been reached.



**model for a DNA molecule at the overstretching transition**

**Figure 2.** Top: Overstretching transition of double-stranded DNA. The force extension relation exhibits a rapid worm-like chain increase until at around 65 pN a plateau is reached. The transition on the plateau corresponds to progressive mechanical denaturation from dsDNA (left end) to ssDNA (right end). Bottom: Schematic depiction of DNA conformation on the overstretching plateau with local denaturation zones (bubbles). See text for details. Figure courtesy Mark C. Williams.

While the double helix is the thermodynamically stable configuration of the DNA molecule below the melting temperature  $T_m$  (at non-denaturing pH), even at physiological conditions there exist local denaturation zones, so-called DNA-bubbles, predominantly in A-T-rich regions of the genome. Driven by ambient thermal fluctuations, a DNA bubble is a dynamical entity whose size varies by thermally activated zipping and unzipping of successive base pairs at the two forks where the ssDNA bubble is bordered by the dsDNA-helix. This incessant zipping and unzipping leads to a random walk in the bubble-size coordinate, and to a finite lifetime of DNA-bubbles under non-melting conditions, as eventually the bubble closes due to the energetic preference for the closed state. This so-called DNA breathing typically opens up a few bps. It has been demonstrated recently that by fluorescence correlation methods the fluctuations of DNA bubbles can be explored on the single molecule level, revealing a multistate kinetics that corresponds to the picture of successive zipping and unzipping of single base pairs. At room temperature, the characteristic closing time of an unbounded base pair was found to be in the range 10 to 100  $\mu\text{sec}$  corresponding to an overall bubble life time in the range of a few msec. The multistate nature of the DNA-breathing was recently confirmed by a UV-light absorption study.

The presence of fluctuating DNA-bubbles is essential to the understanding of the binding of single-stranded DNA binding proteins (SSBs) that selectively bind to ssDNA and that play important roles in replication, recombination and repair of DNA. One of the key tasks of SSBs is to prevent the formation of secondary structure in ssDNA. From the thermodynamical point of view one would therefore expect SSBs to be of an effectively helix-destabilizing nature, and thus to lower  $T_m$ . However, it was found that neither the gp32 protein from the T4 phage nor *E.coli* SSBs do. An explanation to this apparent paradox was suggested to consist in a kinetic block, i.e., a kinetic regulation such that the rate constant for the binding of SSBs is smaller than the one for bubble closing. This hypothesis could recently be verified in extensive single molecule setups using mechanical overstretching of dsDNA by optical tweezers in the presence of T4 gene 32 protein [1,2]. Our study consisted in developing a biophysical model of this system, as detailed below; see references [3-7], and references cited therein.

## B. Probabilistic modeling by the master equation (ME)

DNA breathing in presence of SSBs is described by the probability distribution  $P(m, n, t)$  to find at time  $t$  a bubble with  $m$  open bps and with  $n$  SSB units bound to the two arches of the bubble. The time evolution of  $P(m, n, t)$  is given by a master equation

$$\frac{\partial}{\partial t} P(m, n, t) = W P(m, n, t) \quad (2)$$

where the transfer matrix  $W$  incorporates rates  $t^\pm$  for bubble increase / decrease as well as rates  $r^\pm$  for SSB binding / unbinding:

$$\begin{aligned}
\frac{\partial}{\partial t} P(m, n, t) = & t^+(m-1, n)P(m-1, n, t) + t^-(m+1, n)P(m+1, n, t) \\
& - \{t^+(m, n) + t^-(m, n)\}P(m, n, t) \\
& + r^+(m, n-1)P(m, n-1, t) + r^-(m, n+1)P(m, n+1, t) \\
& - \{r^+(m, n) + r^-(m, n)\}P(m, n, t)
\end{aligned} \tag{3}$$

The rates  $t^\pm$  and  $r^\pm$  are based on the statistical weight  $Z(m, n)$  to find a state  $(m, n)$  and fulfill the detailed balance condition

$$\begin{aligned}
t^+(m-1, n)Z(m-1, n) &= t^-(m, n)Z(m, n) \\
t^+(m, n-1)Z(m, n-1) &= t^-(m, n)Z(m, n)
\end{aligned} \tag{4}$$

$Z(m, n)$  has two contributions,

$$Z(m, n) = Z_{\text{bubble}}(m)Z_{\text{SSB}}(m, n) \quad . \tag{5}$$

The bubble part, according to the Poland-Scheraga model for DNA-melting, is given by

$$Z_{\text{bubble}}(m) = \sigma_0 u^m (m+1)^{-c} \quad , \quad m \geq 1 \quad , \tag{6}$$

with the bubble initiation factor  $\sigma_0$ , the weight  $u$  for breaking a base-pair, and the loop closure factor  $(m+1)^{-c}$  for creating a polymer loop of size  $m$ , with the offset by 1 due to persistence length corrections. We choose typical values,  $\sigma_0 = 10^{-3}$  and  $c = 1.76$ .

Equation (5) is completed by the boundary condition  $Z_{\text{bubble}}(0) = 1$ . The contribution from the SSBs has the form

$$Z_{\text{SSB}}(m, n) = \kappa^n \Omega(m, n) \tag{7}$$

with the binding strength  $\kappa = c_0 K^{\text{eq}}$ , involving the SSB concentration  $c_0$  and equilibrium binding constant  $K^{\text{eq}} = v_0 e^{\beta|E_{\text{SSB}}|}$ , where  $v_0$  is the typical SSB volume and  $E_{\text{SSB}}$  its binding energy. The weight  $\Omega(m, n)$  is a combinatorial factor, counting all possible ways of putting  $n$  SSBs onto the two arches of the bubble, taking into account gaps between SSBs due to the fact that their size  $\lambda$  is larger than a base. Solving the above equations results in the rates for the bubble size  $t^\pm$ :

$$\begin{aligned}
t^+(m,n) &= k u \left( \frac{m+1}{m+2} \right)^c, \quad m \geq 1 \\
t^-(m,n) &= k \frac{\Omega(m-1,n)}{\Omega(m,n)}
\end{aligned} \tag{8}$$

with the bubble initiation rate  $t^+(0,0) = 2^{-c} k \sigma_0 u$ . The SSB number transfer rates  $r^\pm$  result as

$$\begin{aligned}
r^+(m,n) &= \gamma k \kappa (n+1) \frac{\Omega(m,n+1)}{\Omega(m,n)} \\
r^-(m,n) &= n \gamma k
\end{aligned} \tag{9}$$

with the dimensionless ratio  $\gamma = q/k$  of the SSB unbinding rate  $q$  and the base pair zipping rate  $k$ .

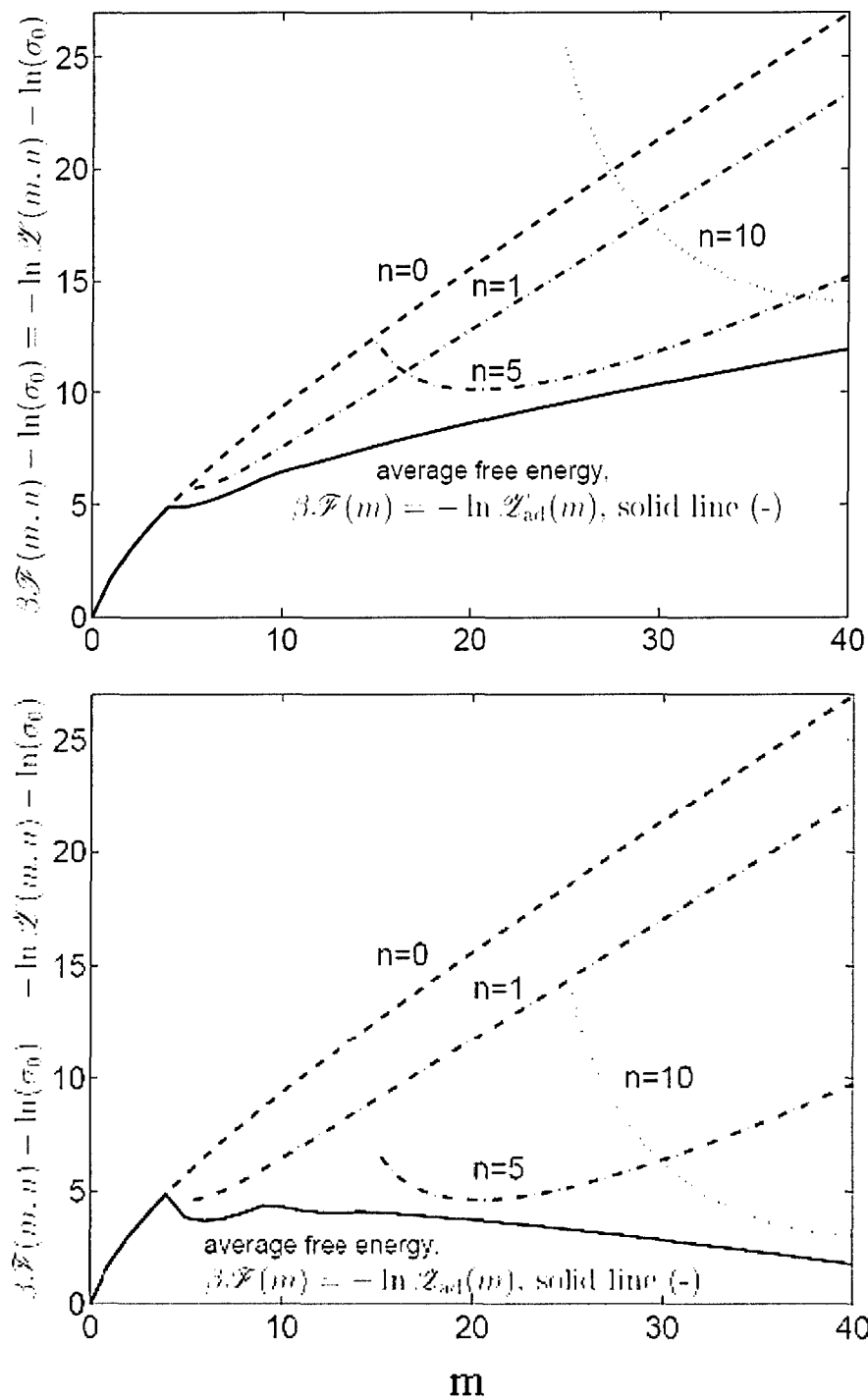
The master equation (3) can be solved by means of an eigenmode expansion of the form

$$P(m,n,t) = \sum_p c_p Q_p(m,n) \exp(t/\tau_p) \tag{10}$$

in which the coefficients  $c_p$  of a given eigenmode  $p$  are determined via the initial conditions. The corresponding eigenvalue equation for the bubble size-SSB number eigenfunction  $Q_p$  determines the mode relaxation times  $\tau_p$  and can be solved numerically.

### C. Effective binding free energy in the limit of fast SSB binding

Fast SSB-binding corresponds to the limit in which the dimensionless ratio  $\gamma = q/k$  of the SSB unbinding rate  $q$  and the base pair zipping rate  $k$  is large, i.e.,  $\gamma \gg 1$ . This limit allows one to average out the SSB-dynamics and to calculate an effective free energy, in which the bubble dynamics with the slow variable  $m$  runs off. The result for two different binding strengths  $\kappa$  is shown in **Figure 3**, along with the free energies corresponding to keeping  $n$  fixed. It is distinct that while for lower  $\kappa$  the presence of SSBs diminishes the slope of the effective free energy, for larger  $\kappa$  the slope becomes negative. This implies that in the first case the bubble opening is more likely, but still globally unfavorable. In the latter case, the presence of SSBs indeed leads to full denaturation. One observes distinct finite size effects due to  $\lambda > 1$ : only when the bubble reaches a minimal size  $m \geq \lambda$ , SSB-binding may occur, a second SSB is allowed to bind to the same arch only once  $m \geq 2\lambda$ , etc. This effect also produces the nucleation barrier for full denaturation in the lower plot of **Figure 3**.



**Figure 3.** Effective free energy in the limit  $\gamma \rightarrow 1$  (solid line), and free energy for various fixed  $n$  ( $u = 0.6$ ,  $c = 1.76$ ,  $\lambda = 5$ ). Top:  $\kappa = 0.5$ ; bottom:  $\kappa = 1.5$ .

#### D. References

- [1] K. Pant, R. L. Karpel, I. Rouzina, and M. C. Williams, J. Mol. Biol. **336**, 851 (2004).
- [2] K. Pant, R. L. Karpel, I. Rouzina, and M. C. Williams, J. Mol. Biol. **349**, 317 (2005).
- [3] A. Hanke and R. Metzler, Phys. Rev. Lett. **90**, 159801 (2003).
- [4] A. Hanke and R. Metzler, J. Phys. A **36**, L473 (2003).
- [5] T. Ambjörnsson and R. Metzler, J. Phys. Cond. Mat. **17**, S1841 (2005).
- [6] T. Ambjörnsson and R. Metzler, Phys. Rev. E **72**, 030901(R) (2005).
- [7] R. Metzler, T. Ambjörnsson, A. Hanke, Y. Zhang, and S. D. Levene, Journal of Computational and Theoretical Nanoscience (in press, 2006).  
E-print physics/0609139.